## AMENDMENTS TO THE CLAIMS:

This listing of the claims will replace all prior versions, and listings of cliams in the application:

## Listing of Claims

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- 20. (Currently Amended) A non-naturally occurring mutant human hemoglobin wherein the leucine residue at position 105 of the  $\beta$ -chains (SEQ ID NO:-7) (SEQ ID.NO:8) is replaced by a tryptophan residue.
- 21. (Original) The hemoglobin of Claim 20 possessing low oxygen affinity as compared to normal human adult hemoglobin.
- 22. (Original) The hemoglobin of Claim 21 further possessing high cooperativity in oxygen binding comparable to normal human adult hemoglobin.
- 23. (Original) The hemoglobin of Claim 20 which is produced recombinantly.
- 24. (Currently Amended) rHb (βL105W) (SEQ ID NO:7)
  (SEQ ID NO:8).

- 25. (Currently amended) An artificial mutant hemoglobin which in a cell-free enviornment has oxygen binding properties comparable to those of human normal adult hemoglobin in red blood cells wherein said hemoglobin contains a mutation such that the leucine residue at position 105 of the  $\beta$ -chains is tryptophan (SEQ ID NO: 7) (SEQ ID NO: 8).
- 26. (Original) The hemoglobin of Claim 25 which is produced recombinantly.
- 27. (Currently Amended) A non-toxic pharmaceutical composition comprising a non-naturally occurring mutant hemoglobin wherein the leucine residue at position 105 of the  $\beta$ -chains is replaced by a tryptophan residue (SEQ ID NO:7) (SEQ ID NO:8) in a pharmaceutically acceptable carrier.
- 28. (Original) The composition of Claim 27 wherein said hemoglobin in a cell-free environment has oxygen binding properties lower than those of human normal adult hemoglobin.
- 29. (Currently Amended) The composition of Claim 28 wherein said hemoglobin is rHb ( $\beta$ L105W) (SEQ ID NO:7) (SEQ ID NO:8).

- 31. (Currently Amended) A non-naturally occurring low oxygen affinity mutant hemoglobin that has oxygen binding properties comparable to those of human normal adult hemoglobin in the presence of the allosteric effector 2,3-bisphosphoglycerate, wherein the leucine residue at position 105 of each of the β-chains is replaced by a tryptophan residue (SEQ ID NO:8).
- mutant human hemoglobin wherein said hemoglobin contains a mutation of the leucine residue at position 105 of the  $\beta$ -chains (SEQ ID NO:7) possessing is replaced by a tryptophan residue (SEQ ID NO:8), wherein said hemoglobin possesses oxygen-binding properties of oxygen affinity as measured by  $P_{50}$  and cooperativity as measured by the Hill coefficient ( $n_{max}$ ) and similar to those of Hb A in the presence of the allosteric effector 2,3-bisphosphoglycerate as follows:  $P_{50}$  about 28.2 mm Hg,  $n_{max}$  about  $\frac{2.6}{2.60}$  in 0.1 M sodium phosphate at pH 7.4 and  $29^{\circ}\text{C}$ .
- 36. (Currently Amended) A method of treating a human subject, comprising administering to said subject a nontoxic composition comprising an artificial mutant hemoglobin, wherein said artificial mutant hemoglobin is rHb ( $\beta$ L105W) (SEQ ID NO:7) (SEQ ID NO:8).

- 37. (Currently Amended) rHb (βL105W) (SEQ ID NO:8). derived from cells transformed with pHE7004.
- 38. (New) The hemoglobin of Claim 22 further possessing increased stability against autooxidation.